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(54) **L-lysyl-glycyl-L-histidine and therapeutic agent for wound healing containing the same**

L-Lysyl-Glycyl-L-Histidin und entsprechende pharmazeutische Zusammensetzungen zur Wundheilung

L-lysyl-glycyl-l-histamine et composition pharmaceutique le contenant pour la cicatrisation de lésions

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(73) Proprietor: **HOECHST JAPAN LIMITED
Tokyo (JP)**

(72) Inventors:
• **Koyama, Masayoshi
Kawagoe-shi, Saitama 350-11 (JP)**
• **Takahashi, Mikiko
Hasuda-shi, Saitama 349-01 (JP)**

• **Yanagawa, Masayoshi
Sayama-shi, Saitama 350-13 (JP)**

(74) Representative: **Lauppe, Hans Friedrich, Dr.
Behringwerke AG
Abt. Patente & Verträge,
Postfach 1140
35001 Marburg (DE)**

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copper-binding peptides that stimulate
angiogenesis'**

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D scriptionBackground of the Invention

5 1. Field of the Invention

[0001] This invention relates to a new tripeptide having an amino acid sequence L-lysyl-glycyl-L-histidine and/or metal complexes thereof, and a pharmaceutical composition for wound healing containing the same as an active ingredient.

10 2. Description of the Prior Art

[0002] Various growth factors are known to be involved in the process of wound healing (Dijke et al., Biotechnology, Vol. 7, p. 793-798, 1989). TGF- β (transforming growth factor- β) and PDGF (platelet-derived growth factor), in particular,

are known to proliferate fibroblasts, induce the cells to a damaged site and enhance wound repairing. Some tripeptides and metal complexes thereof have been known to have a cell proliferation effect. For example, glycyl-L-histidyl-L-lysine, which was isolated from plasma or synthesized and metal complexes thereof, such as, glycyl-L-histidyl-L-lysine : copper (II) are involved in cell proliferation of hepatocytes, neurocytes, kidney cells, thyroid cells, etc., and maintenance of growth (Picart, L. Method, Enzym. Vol. 147, p. 314-328, 1987; Picart, L. IN Vitro, Vol. 17, p. 459-466, 1981; Picart, L. US Patent No. 4,760,051). It is assumed that the cell proliferation activity of glycyl-L-histidyl-L-lysine involves binding of an essential metal ion of trace elements such as copper ion and incorporating the copper ion in cells (Picart, L. IN Vitro, Vol. 17, p. 459-466, 1981). Also disclosed is a use of glycyl-L-histidyl-L-lysine and metal complexes thereof, for example, glycyl-L-histidyl-L-lysine : copper (II) as an agent for promoting a therapy of affected bones and other hard tissues such as cartilage of warm-blooded animals (Picart, L. R., Japanese Patent Publication No. Hei 5-501253).

WO91/03488 discloses glycyl-L-histidyl-L-lysine and L-lysyl-L-histidyl-glycine and copper (II) salts thereof for treating ulcers.

L-lysyl-L-histidyl-glycine was isolated from an avian bursa fabricii as a B cell differentiating hormone (Audhya, T. et al. Science Vol. 231, p. 997-999, 1986). Also it is disclosed that L-lysyl-L-histidyl-glycine is used as an agent for promoting a therapy for the affected bones and other hard tissues such as cartilage (Picart, L. R., the above-mentioned Patent Publication).

L-histidyl-glycyl-L-lysine and L-histidyl-L-lysyl-glycine have been isolated from the feline spinal marrow as an inhibitor of transmission of neurocytes (Lote, C.J. et al., Nature, Vol. 264, p. 188-189, 1976). On the other hand, only glycyl-L-histidyl-L-lysine is known to constitute a metal complex with a metal ion, for example, copper ion among the above-mentioned tripeptides (Picart, L., the afore-mentioned).

However, it has never been reported that L-lysyl-glycyl-L-histidine nor metal complexes thereof, for example, L-lysyl-glycyl-L-histidine : copper (II) proliferates fibroblasts or is effective for wound healing.

Detailed Description of the Invention

[0003] Consequently, an object of the invention is to provide a tripeptide having a wound healing activity.

[0004] Another object of the invention is to provide a pharmaceutical composition for wound healing containing said tripeptide as an effective ingredient.

[0005] TGF- β and PDGF, which were hitherto considered to have a wound healing effect, are physiologically active substances having various functions of themselves and are considered to have other undesirable activities than wound healing. Therefore, a low molecular peptide having no such side effects and being specifically active in wound healing has been desired for the above purposes.

[0006] The present inventors have found, as a result of various studies, the fact that L-lysyl-glycyl-L-histidine and/or metal complexes thereof such as L-lysyl-glycyl-L-histidine : copper (II) promotes fibroblast proliferation, and this invention has now been completed.

[0007] The tripeptide and metal complexes thereof according to this invention are highly water-soluble, and they are most suitably administered in combination with a suitable water-soluble base for wound healing preferably by applying locally to an affected region. Preparations for an external use according to the invention may take the form of a water-soluble ointment, an oleaginous ointment, lotion, spray, oil, gel or the like. Representative bases may include macrogols for a water-soluble ointment, vaselin for an oleaginous ointment, vegetable oils such as olive oil, sesame oil, camellia oil and the like for an oil preparation, and carboxy vinyl polymer, sodium polyacrylate and the like for a gel preparation. The tripeptide and metal complexes thereof according to the invention can also be administered intravenously or subcutaneously in systemic administration, and nasally or transpulmonarily in the form of micronized aerosols.

sols.

[0008] The dosages are in a range of 1 to 100 mg/administration site/person/day in local administration, and 0.1 to 10 mg/kg/day in systemic administration.

[0009] The following examples are provided to illustrate the invention and are not intended to limit the invention.

Examples

Example 1 Synthesis of L-lysyl-glycyl-L-histidine

[0010] A peptide consisting of L-lysyl-glycyl-L-histidine was synthesized by way of solid-phase synthesis using an automatic peptide synthesizer (Applied Biosystems, Inc., U.S.A.). Using 0.5 mM of resin comprising styrene-divinylbenzene copolymer (molar ratio; styrene : divinylbenzene = 99 : 1), amino acids were connected successively toward the N-terminus of the peptide. One mM of N-(t-butoxycarbonyl)-L-Lys, N-(t-butoxycarbonyl)-Gly and N-(t-butoxycarbonyl)-L-His, respectively, were used as amino acids in the reaction. The peptides were obtained by the detachment from the solid-phase and the removal of protecting group by using 5 ml of 95% trifluoroacetic acid (TFA). The peptides obtained were purified by HPLC (available from Hitachi Corp.), then reverse-phase C-18 column (available from Vydac, Inc.) eluting with a linear gradient of acetonitrile containing 0.1% TFA.

Example 2 Determination of fibroblast proliferation promoting activity of L-lysyl-glycyl-L-histidine

[0011] Fibroblast cell strains, Balb/3T3 cells (purchased from ATCC) were inoculated into a 96-well culture plate at 5×10^3 cells/well, 100 μ l of 10% calf serum-containing Dulbecco Modified Eagle Medium (hereinafter referred to as DME) was added and incubated at 37°C for 24 hours in an incubator. Then, the culture medium was removed and the cells were washed. One hundred μ l of lowered serum medium (0.2% calf serum-containing DME) was added to the wells and incubation was continued for another 3 days. L-lysyl-glycyl-L-histidine obtained in Example 1 was added thereto at 10 μ l/well and incubation was performed for 15 hours. The culture medium contains a trace amount of Cu²⁺, and a part of L-lysyl-glycyl-L-histidine may form copper complexes. The 3H-thymidine was added to be 74 KBq/ml and incubation was performed for 6 hours. After completion of the incubation, the medium was removed, the cells were collected and the amount of 3H-thymidine incorporated in the cells was determined.

[0012] The results of the determination of fibroblast proliferation promoting activity of L-lysyl-glycyl-L-histidine are shown in Table 1, wherein the data show the mean and its standard deviation (4 cases per one group).

Table 1

Added Compound	Dose (M)	Incorporated 3H-thymidine (cpm)
Control	-	1392.3 \pm 348.2
L-lysyl-glycyl-L-histidine	10 ⁻⁶	1406.5 \pm 470.5
	10 ⁻⁵	1371.8 \pm 249.1
	10 ⁻⁴	4889.8 \pm 495.1
	10 ⁻³	7719.5 \pm 1246.8

[0013] From the results, it was confirmed that L-lysyl-glycyl-L-histidine has a dose-dependent cell proliferation promoting activity.

Claims

1. A tripeptide comprising an amino acid sequence L-lysyl-glycyl-L-histidine and/or a metal complex thereof.
2. The tripeptide according to claim 1 comprising an amino acid sequence L-lysyl-glycyl-L-histidine.
3. The tripeptide metal complex according to claim 1 comprising L-lysyl-glycyl-L-histidine : copper (II).
4. A pharmaceutical composition for wound healing containing an effective amount of a tripeptide comprising an amino acid sequence L-lysyl-glycyl-L-histidine and/or a metal complex thereof as an active ingredient.

5. The pharmaceutical composition for wound healing according to claim 4 wherein the tripeptide comprises an amino acid sequence L-lysyl-glycyl-L-histidine.
6. The pharmaceutical composition for wound healing according to claim 4 wherein the tripeptide metal complex comprises L-lysyl-glycyl-L-histidine : copper (II).

Patentansprüche

1. Ein Tripeptid, welches eine Aminosäuresequenz L-Lysyl-Glycyl-L-Histidin umfaßt, und/oder ein Metallkomplex hiervon.
2. Das Tripeptid nach Anspruch 1, welches eine Aminosäuresequenz L-Lysyl-Glycyl-L-Histidin umfaßt.
3. Der Tripeptidmetallkomplex nach Anspruch 1, welcher L-Lysyl-Glycyl-L-Histidin:Kupfer(II) umfaßt.
4. Eine pharmazeutische Zusammensetzung zur Wundheilung, welche eine wirksame Menge eines Tripeptids, welches eine Aminosäuresequenz L-Lysyl-Glycyl-L-Histidin umfaßt, und/oder eines Metallkomplexes hiervon als wirksamen Bestandteil enthält.
5. Pharmazeutische Zusammensetzung zur Wundheilung nach Anspruch 4, worin das Tripeptid eine Aminosäuresequenz L-Lysyl-Glycyl-L-Histidin umfaßt.
6. Pharmazeutische Zusammensetzung zur Wundheilung nach Anspruch 4, worin der Tripeptidmetallkomplex L-Lysyl-Glycyl-L-Histidin:Kupfer(II) umfaßt.

Revendications

1. Tripeptide comprenant une séquence d'acides aminés L-lysyl-glycyl-L-histidine et/ou complexe métallique de celui-ci.
2. Tripeptide selon la revendication 1, comprenant la séquence d'acides aminés L-lysyl-glycyl-L-histidine.
3. Complexe métallique de tripeptide selon la revendication 1, comprenant le complexe L-lysyl-glycyl-L-histidine:cuivre-(II).
4. Composition pharmaceutique pour la cicatrisation, contenant une quantité efficace d'un tripeptide comprenant comme composant actif la séquence d'acides aminés L-lysyl-glycyl-L-histidine et/ou d'un complexe métallique de celui-ci.
5. Composition pharmaceutique pour la cicatrisation selon la revendication 4, dans laquelle le tripeptide comprend la séquence d'acides aminés L-lysyl-glycyl-L-histidine.
6. Composition pharmaceutique pour la cicatrisation selon la revendication 4, dans laquelle le complexe métallique de tripeptide comprend le complexe L-lysyl-glycyl-L-histidine:cuivre-(II).